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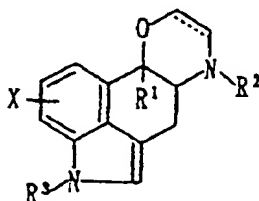
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D-80469 München (DE)(54) **Tetracyclic morpholine derivatives and their use or analgesics.**

(57) A compound represented by the formula



wherein X is hydrogen or a substituent; R¹ is an optionally substituted hydrocarbon residue; R² and R³ are respectively hydrogen or a substituent; and ----- shows a single bond or a double bond, or a pharmacologically acceptable acid addition salt thereof, has an analgesic action and is useful as a medicine.

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This invention relates to novel condensed morpholine derivatives.

Compounds having an indolobenzoxazine skeleton were disclosed in USP-4,493,836 and USP-4,318,910 by Lucien Nedelec et al., and in USP-4,238,486 by James H. Jones, in which anti-Parkinson and antihypertensive activity are indicated. In these known compounds, the carbon atom at 1a position of the indolobenzoxazine skeleton is unsubstituted.

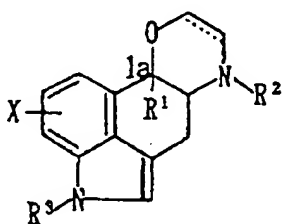
Pain is a complaint due to physical or mental causes, and man has endeavored, since ancient times, to synthesize drugs capable of eliminating or relieving the pain.

Stimuli causing pain include, among others, chemical, electrical, thermal and mechanical ones. Pain signals given to primary sensory nerve by these stimuli are transmitted to cerebral cortex through spinal cord, where the sensation is recorded as pain. In the transmission of pain, various neuropeptides are participated. Above all, substance-P (SP) is a peptide consisting of 11 amino acids belonging to tachykinins, which is counted as one of the substances of transmitting information from primary sensory nerve to the dorsal horn of the spinal cord, which participates in, other than pain, central nervous system diseases, inflammatory diseases. Therefore, drugs capable of suppressing the stimuli transmitting the action of SP can be used as excellent analgesics.

As a strong analgesic which has been known best, mention is made of a narcotic analgesic, e.g. morphine, and a number of its related compounds have been synthesized. Most of these drugs, however, cause tolerance and physical and mental dependency, and the strength of such tolerance and dependency is assumed to be in direct proportion to the analgesic activity. Circumstances being such as above, development of non-narcotic analgesic which does not act on a morphine-acceptor has been ardently desired.

The present inventors found that novel indolobenzoxazine derivatives modified by introducing into their 1a-position a hydrocarbon residue such as lower alkyl, lower alkenyl, lower alkynyl or aryl presented analgesic actions in various animal models, and then conducted further investigation on the finding to complete the present invention.

Thus, this invention relates to: a compound (I) represented by the formula



(Formula 1)

wherein X is hydrogen or a substituent; R¹ is an optionally substituted hydrocarbon residue; R² and R³ are respectively hydrogen or a substituent; and — shows a single bond or a double bond, or a pharmacologically acceptable acid addition salt thereof, and the use of the compound as an analgesics.

Referring to the compound (I) above, more particularly, hydrogen, halogen or an optionally substituted hydrocarbon residue which may be bound to the ring through oxygen atom are mentioned for X;

an optionally substituted hydrocarbon residue is mentioned for R¹;

hydrogen or an optionally substituted hydrocarbon residue are mentioned for R²;

hydrogen, an optionally substituted hydrocarbon residue or a sulfonyl group are mentioned for R³.

More preferably, X is hydrogen, a lower alkyl, a lower alkoxy or halogen;

R¹ is a lower alkyl, a lower alkenyl, a lower alkynyl or aryl;

R² is hydrogen or an optionally substituted lower alkyl;

R³ is hydrogen or an optionally substituted lower alkyl or sulfonyl group.

Referring to the above-mentioned X, R¹, R² and R³, fluorine, chlorine, bromine and iodine are mentioned as halogen. The lower alkyl includes C₁₋₆ straight-chain or branched saturated hydrocarbon, one or more hydrogen atoms of which being optionally substituted with halogen, as exemplified by methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, n-pentyl, n-hexyl, trifluoromethyl, difluoromethyl, monofluoromethyl, pentafluoroethyl and 2-trifluoroethyl.

The lower alkenyl includes C₂₋₆ straight-chain or branched unsaturated hydrocarbon having a double bond in the molecule, as exemplified by vinyl, allyl, 2-propenyl, 2-butenyl, isoprenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl and 5-hexenyl.

The lower alkynyl includes C₂₋₆ straight-chain or branched unsaturated hydrocarbon having a triple bond in the molecule, as exemplified by propargyl, ethynyl, 2-propynyl, 3-propynyl, 1-hexynyl, 2-hexynyl, 3-

hexynyl and 4-hexynyl.

The lower alkoxy includes functional groups formed by binding the above-mentioned lower alkyl through oxygen atom, as exemplified by C₁₋₆ alkoxy such as methoxy, ethoxy, n-propoxy, n-butoxy, t-butoxy, isopropoxy and trifluoromethoxy.

5 The aryl includes monocyclic or condensed polycyclic aromatic hydrocarbon, as exemplified by phenyl, naphthyl, indenyl and anthryl.

As substituents of the optionally substituted lower alkyl group shown by the above-mentioned R² and R³, mention is made of, for example, the above-mentioned halogen, aryl or cycloalkyl, heterocyclic ring, aralkyl and acyl, and the aryl and heterocyclic ring may optionally be bound to the lower alkyl through
10 oxygen, sulfur or nitrogen atom.

The cycloalkyl includes 3- to 7-membered ring formed with solely carbon atoms, which may optionally have a double bond in the ring, as exemplified by cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, 2-cyclohexen-1-yl and 2-cyclopenten-1-yl.

The heterocyclic ring includes rings which may be 4- to 7-membered monocyclic or condensed cyclic
15 rings optionally having unsaturated bond and having at least one of oxygen atom, nitrogen atom, and sulfur atom as the ring-constituting atom, as exemplified by 2-thienyl, 3-thienyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 2-furyl, 3-furyl, 4-quinolyl, 8-quinolyl, 4-isoquinolyl, pyrazinyl, 2-pyrimidinyl, 1-pyrrolidinyl, 2-pyrrolidinyl, 3-pyrrolidinyl, 2-imidazolyl, 3-imidazolyl, 1-piperidyl, 3-piperidyl, 1-morpholinyl, 2-morpholinyl, 3-morpholinyl, 4-morpholinyl, 1-piperazinyl, 2-piperazinyl, 3-piperazinyl, 2-isindolinyl and 1-phthalimide.

20 The aralkyl includes functional groups formed by binding one or more aryl groups through lower alkyl, as exemplified by benzyl, phenethyl, diphenylmethyl, triphenylmethyl, 1-naphthylmethyl, 2-naphthylmethyl, 2-diphenylethyl, 3-phenylpropyl, 4-phenylbutyl and 5-phenylpentyl.

The acyl includes functional groups formed by binding the above-mentioned lower alkyl, lower alkenyl, lower alkynyl, aryl, cycloalkyl, aralkyl or heterocyclic ring through carbonyl, phosphoryl or sulfonyl, as
25 exemplified by formyl, acetyl, propionyl, butyryl, valeryl, acryloyl, propioly, benzoyl, nicotinoyl, methanesulfonyl, ethanesulfonyl, phenylsulfonyl, tolylsulfonyl and 2,4,6-triisopropylphenylsulfonyl.

The above-mentioned aryl, aralkyl and heterocyclic ring may optionally have, at optional positions, one or plural optional substituents which are different from one another.

The optional substituents include the above-mentioned lower alkyl, lower alkenyl, lower alkynyl,
30 cycloalkyl or cyano, nitro, sulfonyl, amino, lower monoalkylamino, lower dialkylamino, arylamino, hydroxyl group, lower alkoxy, halogen, carbamoyl, carboxyl, lower alkoxycarbonyl, lower alkanoyl, mercapto, lower alkylmercapto, aralkyl, aryl, lower alkyl carbamoyl, aryl carbamoyl, acyl (e.g. benzoyl, nicotinoyl), methylenedioxy, heterocyclic ring (preferably 4-pyridyl, 2-pyridyl, 3-pyridyl and 2-pyrimidinyl), or the like.

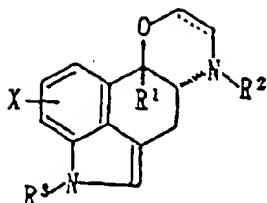
The sulfonyl group for R³ includes a sulfonyl group optionally substituted by lower alkyl or aryl which
35 may be substituted by lower alkyl, preferably arylsulfonyl which may be substituted by lower alkyl, more preferably phenylsulfonyl which may be substituted by one to three C₁₋₆ alkyl groups, as exemplified by p-tolylsulfonyl and 2,4,6-triisopropylphenylsulfonyl.

Among the compounds (I), preferable ones are those wherein X stands for hydrogen or lower alkoxy; R¹ stands for a lower alkyl, a lower alkenyl or phenyl; R² stands for hydrogen, a lower alkyl or a group (II)
40 shown by the formula, -(CH₂)_nAR⁴ (Formula 2): wherein n is an integer of 1 to 6, A is a bond or oxygen, sulfur or nitrogen atom, R⁴ is hydrogen, phenyl, 1-phthalimidyl, 4-benzylpiperazyl; R³ stands for hydrogen, 2,4,6-triisopropylphenylsulfonyl or p-tolylsulfonyl and ----- is a single bond.

More preferable ones are those wherein X is hydrogen or C₁₋₃ alkoxy; R¹ is C₁₋₃ alkyl, vinyl or phenyl; R² is hydrogen, C₁₋₆ alkyl or the group of the formula -(CH₂)_nAR⁴ wherein n is an integer of 1 to 6, A is a
45 bond or oxygen, sulfur or nitrogen atom, R⁴ is hydrogen, phenyl, 1-phthalimidyl, 4-benzylpiperazyl; R³ is hydrogen or 2,4,6-triisopropylphenylsulfonyl and ----- is a single bond.

Examples of pharmacologically acceptable acid addition salts include inorganic salts such as hydrochloride, hydrobromide, hydroiodide, sulfate or phosphate, and organic salts such as acetate, oxalate, succinate, ascorbate, maleate, lactate, citrate, fumarate, tartrate, methanesulfonate or benzoate.

50 It is preferable that the compound (I) of this invention has the following steric configuration;

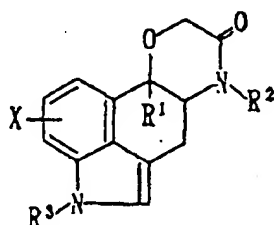


(Formula 3)

wherein X, R¹, R² and R³ are of the same meaning as defined above.

Further, in the compounds included in the present invention, there exist optical isomers, and the optically active compounds obtained by optical resolution of those isomers are also included in the present invention.

The compound (I) of this invention can be produced by, for example, subjecting a compound (IV) shown by the formula



(Formula 4)

wherein X, R¹, R² and R³ are of the same meaning as defined above to reduction, and, when desired, by further subjecting the reduced compound to reduction, oxidation or/and substitution reaction.

The above-mentioned reduction is conducted in an inert solvent by reacting with a metal hydride (e.g. lithium aluminum hydride, lithium boron hydride, sodium cyano borohydride or diborane). As the inert solvent, mention is made of ether (e.g. ethyl ether, tetrahydrofuran or dioxane), halogenated hydrocarbon (chloroform, dichloromethane), hexane, benzene, toluene or the like. The metal hydride is used in a amount of 1 to 20 equivalents, preferably 3 to 12 equivalents, and the reaction temperature ranges from -70 °C to 50 °C. Preferably, the reaction is conducted in an ether solvent at temperatures ranging from 0 °C to 30 °C with lithium aluminum hydride.

Further, introduction and modification of substituents can be conducted, for example, in the following manner.

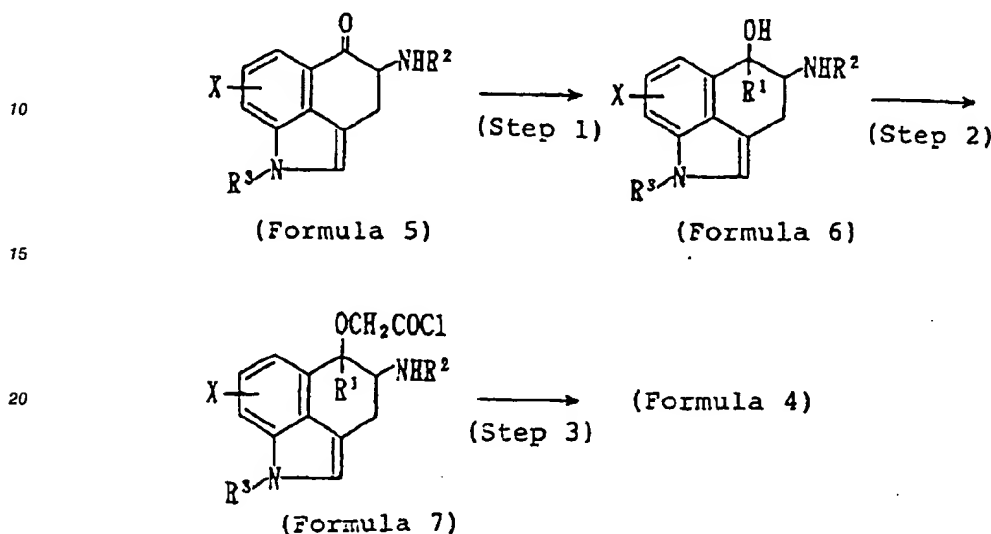
The alkylation on the secondary nitrogen atom is conducted processing with a base (1 to 5 equivalents, preferably 2 to 3 equivalents) in an inert solvent, for example, water, ether (e.g. tetrahydrofuran, ethyl ether or dioxane), acetonitrile, N,N-dimethylformamide and halogenated hydrocarbon, singly or a mixture of them. Examples of the base include potassium carbonate, sodium carbonate, sodium hydride, potassium hydride, sodium ethoxide and potassium-t-butoxide, and the reaction temperature ranges from -50 °C to 80 °C, preferably from 0 °C to 30 °C.

The acylation on the secondary nitrogen atom is conducted by allowing a reactive derivative of chlorine, acetic acid or bromoacetic acid (e.g. chloroacetyl chloride, bromoacetyl bromide or bromoacetyl chloride) to react in an inert solvent. Preferable examples of the inert solvent include water, ether, halogenated hydrocarbon, acetonitrile and N,N-dimethylformamide, and they are used singly or as a mixture. Depending on cases, a base is added to the reaction system, as exemplified by inorganic bases (sodium carbonate, sodium hydrogencarbonate, sodium hydroxide, potassium carbonate, etc.) or organic bases (triethylamine, pyridine, 4,4-dimethylamino pyridine, diisopropyl ethylamine, etc.). The amount of the base ranges from 1 to 10 equivalents, preferably from 1 to 3 equivalents. The reaction temperature ranges from 0 °C to 50 °C, preferably from 10 °C to 30 °C.

The elimination of a base shown by R³ (e.g. sulfonyl derivatives) can be conducted by basic hydrolysis, acid hydrolysis or reductive elimination, and the last one is preferable. The reaction is conducted by processing with sodium naphthalene prepared from metallic sodium and naphthalene or a metal hydride (e.g. lithium aluminum hydride) in an inert solvent (e.g. ether, toluene or hexane, singly or as a mixture). The amount of the reducing agent ranges desirably from 5 to 20 equivalents.

As the inert solvent, tetrahydrofuran is preferably employed, and the reaction temperature ranges from 0°C to 40°C, preferably around room temperature.

The starting compound (Formula 4) can be produced from 3,4-dihydrobenzo[cd]indol-5(1H)-one (J. Chem. Soc., p.1438, 1973) or a derivative thereof by the following steps in accordance with the method described by Bowman (J. Chem. Soc., p.1121, 1972).



wherein X, R¹, R² and R³ are of the same meaning as defined above.

Step 1

This step is a nucleophilic addition reaction on ketone group. The reaction is conducted by reacting with an organometallic reagent represented by R¹-M (M stands for Li, Na, MgBr, MgCl or CeCl₂) in an inert solvent. As the inert solvent, use is made of tetrahydrofuran, ethyl ether, dimethoxyethane or dioxane, singly or as a mixture. The amount of the organometal ranges from 1 to 20 equivalents, preferably from 4 to 10 equivalents. The reaction temperature ranges from -100°C to 70°C, preferably from -20°C to 25°C.

Step 2

This is chloroacetylation reaction, which is conducted in accordance with the method of Lucien Nedelec et al. (JP-A-133292/1981). Namely, the reaction with chloroacetyl chloride (preferably 1 to 2 equivalents) is conducted in an inert solvent in the presence of an excess amount of a base. As the inert solvent, use is made of water, ether (ethyl ether, tetrahydrofuran or dioxane), halogenated hydrocarbon (chloroform, dichloromethane or dichloroethane), ethyl acetate, acetonitrile and N,N-dimethylformamide, singly or as a mixture. The reaction is conducted at temperatures ranging from 0°C to 50°C. Preferable bases are exemplified by potassium carbonate, sodium carbonate, sodium hydroxide and potassium hydroxide, and, preferable inert solvents include tetrahydrofuran, chloroform and dichloromethane. Preferable reaction temperatures range from 0°C to 30°C.

Step 3

This is intramolecular cyclization. The reaction is conducted by processing with a base (1 to 5 equivalents, preferably 2 to 3 equivalents) in an inert solvent, for example, ether (e.g. tetrahydrofuran, ethyl ether or dioxane), acetonitrile, N,N-dimethylformamide or halogenated hydrocarbon, singly or as a mixture. Preferable examples of the base include inorganic strong bases (e.g. sodium hydride, potassium hydride, sodium hydroxide, potassium hydroxide, sodium ethoxide or potassium-t-butoxide), especially preferable one being sodium hydride. Preferable examples of the inert solvent include tetrahydrofuran, acetonitrile and N,N-dimethylformamide, which can be used singly or as a mixture. The reaction temperatures range preferably from 0°C to 40°C.

Optically active compounds can be produced by per se known means, more specifically stating, by using an optically active synthetic intermediate or by subjecting the mixture of the ultimate racemic compounds to optical resolution.

As the optical resolution, mention is made of a method which comprises allowing an optically active acid and salt to be formed, followed by separating them by fractional recrystallization; a method which comprises subjecting a racemic compound or a salt thereof to chromatography using a column for separation of optical active compounds (chiral column), for example, ENANTIO-OVM (Toso Co., Ltd.), followed by developing with a solvent such as water, various buffers (e.g. phosphate buffer), alcoholic solvents (e.g. methanol or ethanol), nitrile type solvents (e.g. acetonitrile), hexane or ethyl ether, singly or as a mixture; and a method which comprises subjecting a mixture of racemic compounds to condensation, by a conventional method (e.g. acid chloride method), with, for example, MPTA (α -methoxy- α -(trifluoromethyl)-phenylacetic acid or menthoxyacetic acid to give a mixture of diastereomers of amido-compound, which is subjected to separation and purification by means of fractional recrystallization or a silica gel chromatography, followed by subjecting thus separated and purified product to acid hydrolysis or basic hydrolysis.

The compound (Formula 1) of this invention is antagonistic against symptoms of pain or the like induced by Substance-P, and shows analgesic activity in animals.

The toxicity of the compounds of this invention is low (acute toxicity (LD_{50}), when Compound 2 of Examples is orally administered to rats, is not less than 100 mg/kg) with less undesirable side effects. Therefore, the compounds of this invention and their salts can be used for prophylaxis or therapy of diseases including pain due to bone diseases (arthritis, rheumatic fever, osteoporosis), pain induced by cancer of the like, lumbago, postoperative pain, neuralgia, pain induced by inflammatory diseases, pain caused by tooth extraction, tooth-ache and pain due to scalds trauma, as well as neurosis (e.g. anxiety neurosis, depression, mental disorder), somniphath or the like. Since the compounds of this invention act on a neurotransmitting system such as dopamine or norepinephrine, they can be use also as the prophylactic and therapeutic drugs of circulatory diseases (e.g. hypertension).

The compounds of this invention and their salts can be safely administered orally or non-orally as they are or as pharmaceutical compositions by being admixed with pharmaceutically acceptable carrier, vehicle or diluent, e.g. tablets (including sugar-coated tablets, film-coated tablets), powders, granules, capsules (including soft capsules), liquids, injections, suppositories, sustained-release drugs.

While the dosage of the compounds and their salts of this invention varies with, for example, the subjects, administration routes and diseases to be treated, it ranges conveniently from 0.1 to 500 mg, preferably 1 to 100 mg, in the case of oral administration to a human adult as an analgesic agent.

Examples

In the following, Na_2SO_4 means sodium sulfate, LAH means lithium aluminum hydride, THF means tetrahydrofuran, NH_4Cl means ammonium chloride, DCE means 1,2-dichloroethane, KBr means potassium bromide, DMF means N,N-dimethylformamide and $CDCl_3$ means deuteriochloroform.

Reference Example 1

3-(5-Methoxy-1-p-toluenesulfonylindol-3-yl)propionic acid

In THF(640 ml)-DMF(130 ml) was dissolved ethyl 3-(5-methoxy-1H-indol-3-yl)propionic acid (99.3 g, 0.4 mol). To the solution was added in limited amounts, under ice-cooling, a suspension of 60% sodium hydride in oil (18.4 g, 0.46 mol) in THF(50 ml)-DMF(10 ml). The mixture was stirred at room temperature for one hour, which was again cooled with ice. To the mixture was gradually added p-toluenesulfonyl chloride (83.9 g, 0.44 mmol), followed by stirring for further 3 hours at room temperature. The reaction mixture was poured into an ice-saturated NH_4Cl solution (180 ml). The upper layer was subjected to extraction with ethyl acetate (100 ml + 500 ml). The extract solution was washed with a saturated aqueous saline solution (300 ml x 2), followed by drying over Na_2SO_4 . The residue was purified by means of a silica gel chromatography (eluent: hexane - ethyl acetate (2:1)) and then were added hydrochloric acid (100 ml) and acetic acid (400 ml), and was refluxed for 1 hour. The reaction mixture was concentrated and the resulting crystals were washed with water, dried under reduced pressure and washed with methanol-isopropyl ether (1:3) to obtain the title compound (54.1 g: m.p. 172-174 °C)

Reference Example 2

3,4-Dihydro-6-hydroxy-1-p-toluenesulfonylbenz [cd] indol-5(1H)-one

To a DCE (600 ml) solution of 3-(5-methoxy-1-p-toluenesulfonyl-indol-3-yl)propionic acid (28.9 g, 77 mmol) was added, while stirring under ice-cooling, (COCl)₂ (8.3 ml, 95 mmol), and the mixture was stirred for about 2.5 hours at room temperature. The reaction mixture was concentrated, which was dissolved in DCE (100 ml). The solution was added to a suspension of aluminum chloride powder (33.9 g, 254 mmol) in DCE (430 ml) at 5 °C taking 20 minutes. The mixture was left standing as it was at 5 °C for one hour, then stirred for 2.5 hours at room temperature. To the reaction mixture were added, under ice-cooling, dimethoxyethane (300 ml) and water (300 ml). Resulting precipitates were filtered off, then the organic layer was separated. The aqueous layer was subjected to extraction with chloroform (200 ml). The extract solution was combined with the organic layer, which was washed with an aqueous saline solution. The resultant organic layer was dried (anhydrous sodium sulfate), then the solvent was distilled off under reduced pressure to leave 25 g of crude crystals. The crude crystalline product was subjected to a silica gel column chromatography, eluting with chloroform, to obtain crystals, followed by recrystallization from chloroform-n-hexane to afford the title compound (m.p. 210-211 °C).

¹H-NMR(CDCl₃, 200MHz) δ: 2.36(s,3H), 2.81(t,J=7Hz,2H), 3.13(t,J=7Hz,2H), 6.8-8.0(m,7H) ppm. IR(KBr): 3095, 1643 cm⁻¹.

Anal: Calcd. for C ₁₈ H ₁₅ NO ₄ S;	C 63.33%,	H 4.43%,	N 4.10%
Found;	C 63.22%,	H, 4.39%,	N 4.08%.

Reference Example 3

3,4-Dihydro-6-methoxy-1-p-toluenesulfonylbenz [cd] indol-5 (1H)-one

To a solution of the crude crystals (42 g) obtained in Reference Example 2 in N,N-dimethylformamide (400 ml) were added potassium carbonate (60.7 g, 440 mmol) and methyl iodide (39.7 g, 280 mmol). The mixture was stirred for about one hour at room temperature, to which were further added potassium carbonate (20.0 g, 140 mmol) and methyl iodide (15.0 g, 110 mmol). The mixture was stirred for further one hour at room temperature. To the reaction mixture was then added water (1000 ml), which was subjected to extraction with ethyl acetate (400 ml x 2). The extract solution was washed with an aqueous saline solution and dried (anhydrous Na₂SO₄). The solvent was then distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with n-hexane-ethyl acetate (1:1) to afford the title compound (48%).

¹H-NMR(CDCl₃, 200 MHz) δ: 2.36(s,3H), 2.78(t,J=7Hz,2H), 3.10(t,J=7Hz,2H), 3.96(s,3H), 6.9-8.1(m,7H), ppm.

IR(KBr): 2940, 1682 cm⁻¹. Anal: Calcd. for C₁₉H₁₇NO₄S; C 64.21%, H 4.82%, N 3.94% Found: C 63.54%, H 4.97%, N 3.93%

Reference Example 4

4-Azido-3,4-dihydro-6-methoxy-1-p-toluenesulfonyl benz [cd]indol-5(1H)-one

To a solution of 3,4-dihydro-6-methoxy-1-p-toluene sulfonylbenz[cd]indol-5(1H)-one (5.1 g, 14.3 mmol) in THF (60 ml) was gradually added, under stirring at -45 °C, a solution of Me₃PhNBr₃ (5.38 g, 14.3 mmol) in THF (20 ml), and the mixture was warmed up to 10 °C. After completion of the reaction, insolubles were filtered off. The filtrate was subjected to distillation under reduced pressure to leave 7.5 g(100%) of 3,4-dihydro-4-bromo-1-(p-toluenesulfonyl) benz[cd]indol-5-one as crude crystals. Then, this crude crystalline product was dissolved in N,N-dimethylformamide-acetic acid (140 ml-2 ml). To the solution was gradually added, under stirring at -28 °C, an aqueous solution (14 ml) of sodium azide (1.86 g, 28.6 mmol) to raise the temperature to about 0 °C. After completion of the reaction, the reaction mixture was added to ice-water (300 ml). Resulting crystalline precipitates were collected by filtration and, then dissolved in ethyl acetate, and the solution was shaken to make two layers. The organic layer was washed with an aqueous saline solution and dried (anhydrous Na₂SO₄). The solvent was then distilled off under reduced pressure to leave

5.67 g (100%) of the title compound as crude crystals. A portion of the crystalline product was recrystallized from ethyl acetate to afford the product, m.p.147-148 °C.

¹H-NMR(CDCl₃, 200 MHz) δ: 2.36(s,3H), 3.02(ddd,J=20, 12, 2 Hz, 1H), 3.33(dd,J=20, 7Hz,1H), 4.00(s,3H), 4.35(dd, J=12, 7Hz, 1H), 6.9-8.1(m,7H) ppm. IR(KBr): 2104, 1688 cm⁻¹. Anal: Calcd for C₁₉H₁₆N₄O₄S.0.3H₂O; C 56.78%, H 3.83%, N 13.82% Found; C 57.03%, H 3.91%, N 13.42%.

Reference Example 5

4-Acetylamino-3,4-dihydro-6-methoxy-1-p-toluenesulfonylbenz[cd]indol(1H)-one

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To a solution of 4-azido-3,4-dihydro-6-methoxy-1-p-toluenesulfonyl benz[cd]indol-5(1H)-one (5.36 g, 13.5 mmol) in THF (130 ml) were added acetic anhydride (2.76 g, 27.0 mmol) and 10% palladium-carbon (2.2 g). The mixture was stirred for 10 hours at room temperature under hydrogen atmosphere of 3 atmospheric pressure. After completion of the reaction, the catalyst was filtered off. The filtrate was subjected to distillation under reduced pressure. The residue was washed with ethyl ether to give 5.03 g (90%) of the title compound as a crude crystalline product. A portion of the product was recrystallized from chloroform - n-hexane to afford a pure product (m.p.214-217 °C).

¹H-NMR(CDCl₃, 200 MHz) δ: 2.37(s,3H), 2.73(ddd,J=18, 14, 2, 1H), 3.85(dd,J=18, 10, 1H), 3.96(s,3H), 4.73(ddd, J=14, 10, 6, 1H), 6.78(br d, 1H), 6.9-8.1(m,7H) ppm. IR(KBr): 3340, 1682, 1634 cm⁻¹. Anal: Calcd for C₂₁H₂₀N₂O₅S.1.0H₂O: C 58.59%, N 6.51% Found; C 59.02%, H 4.88%, N 6.42%.

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Reference Example 6

4-Amino-3,4-dihydro-6-methoxy-1-p-toluenesulfonyl benz[cd]indol-5(1H)-one hydrochloride

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To a suspension of 4-azido-3,4-dihydro-6-methoxy-1-p-toluenesulfonyl benz[cd]indol-5(1H)-one (2.0 g, 5.0 mmol) in ethanol (100 mmol) were added conc. hydrochloric acid (2.5 ml) and 10% palladium-carbon (400 mg). The mixture was stirred for 4 hours at room temperature under hydrogen atmosphere of 3 atmospheric pressure. After completion of the reaction, the catalyst was filtered off. The filtrate was subjected to distillation, and the residue was washed with ethyl ether to give 1.6 g (80%) of crude crystals of the title compound.

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¹H-NMR(DMSO-d₆, 200 MHz) δ: 3.12(t,J=10Hz, 1H), 3.89(s,3H), 4.44(m,1H), 7.2-8.1(m,7H), 8.50(br s,2H) ppm.

IR(KBr): 1687 cm⁻¹. Anal: Calcd. for C₁₉H₁₉N₂O₄SCl.3.0H₂O; C 49.51%, H 5.47%, N 6.08% Found: C 49.57%, H 5.22%, N 5.95%.

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Reference Example 7

3,4-Dihydro-6-methoxy-1-p-toluenesulfonyl-4-trifluoroacetylaminobenz[cd]indol-5(1H)-one

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To a suspension of 4-amino-3,4-dihydro-6-methoxy-1-p-toluenesulfonylbenz[cd]indol-5(1H)-one hydrochloride (1.45 g, 3.50 mmol) in dichloromethane (30 ml) were added, while stirring under ice-cooling, anhydrous trifluoroacetic acid (0.61 ml, 4.27 mmol) and triethylamine (1.18 ml, 16.1 mmol). The mixture was stirred for 30 minutes as it was. After completion of the reaction, the reaction mixture was subjected to distillation under reduced pressure. The residue was shaken together with ethyl acetate (40 ml)-1N HCl (30 ml) to allow two layers to be formed. The organic layer was washed with an aqueous saline solution and dried (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure. The residue was recrystallized from ethyl acetate - n-hexane to afford the title compound (1.2 g, 73%, m.p.189-190 °C)

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¹H-NMR(CDCl₃), 200 MHz) δ: 2.84(ddd,J=14,13,2Hz,1H),3.98(s,3H), 4.70(m,1H), 7.0-8.2(m,7H) ppm. IR(KBr): 3355, 1732, 1690 cm⁻¹. Anal: calcd. for C₂₁H₁₇N₂O₅SF₃.0.5H₂O; C 53.05%, H 3.82%, N 5.89% Found; C 52.96%, H 3.97%, N 5.80%.

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Reference Example 8

Methyl 1-(2,4,6-triisopropylphenylsulfonyl)-indole-3-propionate

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Sodium hydride (60% oil 19.03 g 0.476 mol) was washed with n-hexane, which was suspended in DMF (500 ml). To the suspension was gradually added a solution of ethyl indole-3-propionate acid (64.79 g 0.317

mol) in DMF(100 ml). The mixture was stirred for one hour at 40-50 °C, to which was gradually added, under ice-cooling, 2,4,6-triisopropylphenylsulfonyl chloride (115.25 g, 0.38 mol), followed by stirring for one hour at room temperature. To the reaction mixture was added water (150 ml), which was subjected to extraction with n-hexane (500 ml). The extract solution was washed with water (100 ml) and dried over anhydrous magnesium sulfate. The solvent was distilled off under reduced pressure to leave the title compound (152 g).

¹H-NMR(CDCl₃, 200 MHz) δ: 1.09(d, J=7Hz, 12H), 1.24(d, J=7Hz, 6H), 2.69(t, J=7Hz, 2H), 2.90-(hep, J=7Hz, 1H), 3.04(t, J=7Hz, 2H), 3.65(s, 3H), 4.16(hep, J=7Hz, 2H), 7.2-7.6(m, 7H) ppm. IR(KBr): 2960, 1740 cm⁻¹.

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Reference Example 9

1-(2,4,6-Triisopropylphenylsulfonyl)-indole-3-propionic acid

15 In acetic acid (200 ml) was dissolved methyl 1-(2,4,6-triisopropylphenylsulfonyl)-indole propionate (152 g). To the solution was added conc. hydrochloric acid (150 ml), and the mixture was refluxed under heating for two hours. The reaction mixture was concentrated under reduced pressure, which was added to a saturated aqueous solution of sodium hydrogencarbonate to render the pH to 5 - 6, followed by extraction with dichloromethane (1000 ml). The extract solution was washed with water (500 ml) and dried over anhydrous magnesium sulfate. The solvent was distilled off under reduced pressure to leave the title compound (153 g) as an oily product.

¹H-NMR(CDCl₃, 200MHz) δ: 1.09(d, J=7Hz, 12H), 1.24(d, J=7Hz, 6H), 2.76(t, J=7Hz, 2H), 2.90-(hep, J=7Hz, 1H), 3.06(t, J=7Hz, 2H), 4.16(hep, J=7Hz, 2H), 7.2-7.6(m, 7H) ppm. IR(KBr): 2955, 1712 cm⁻¹.

25 Reference Example 10

3,4-Dihydro-1-(2,4,6-triisopropylphenylsulfonyl)-benz[cd]indol-5(1H)-one

To a solution of 1-(2,4,6-triisopropylphenylsulfonyl)-indole-3-propionic acid (76.22 g) in chloroform (500 ml) was added dropwise thionyl chloride (28.9 ml) at room temperature taking about 10 minutes. The mixture was stirred for one hour at 60 °C. After completion of the reaction, the solvent was distilled off under reduced pressure. To the residue was again added chloroform, followed by concentration under reduced pressure. The concentrate was dissolved in dichloroethane (900 ml). To the solution was gradually added, under ice-cooling, crushed aluminum chloride (58.4 g). The mixture was stirred for 30 minutes, which was added to a mixture of 1N hydrochloric acid (1000 ml) and ice (200 g), followed by extraction with chloroform (900 ml). The extract solution was washed with water (900 ml) and dried over anhydrous magnesium sulfate, which was concentrated under reduced pressure. The concentrate was purified by means of a column chromatography (eluted with ethyl acetate : n-hexane = 1:4), followed by recrystallization from ethyl acetate - hexane to afford the title compound (43.86 g, m.p.124-130 °C).

¹H-NMR(CDCl₃, 200MHz) δ: 1.10(d, J=7Hz, 12H), 1.25(d, J=7Hz, 6H), 2.87(t, J=7Hz, 2H), 3.21(t, J=7Hz, 2H), 4.18(hep, J=7Hz, 1H), 7.2-7.4(m, 4H), 7.68(d, J=9Hz, 2H) ppm.

Reference Example 11

45 4-Azido-3,4-dihydro-1-(2,4,6-triisopropylphenylsulfonyl)benz[cd]indol-5(1H)-one

To a solution of 3,4-dihydro-1-(2,4,6-triisopropyl phenylsulfonyl)benz[cd]indol-5(1H)-one (5.1 g, 14.3 mmol.) in THF (60 ml) was gradually added under stirring at -45 °C a solution of Me₃PhBr₃ (5.38 g, 14.3 mmol.) in THF (20 ml) to raise the temperature of the reaction system up to 10 °C. After completion of the reaction, insolubles were filtered off, and the filtrate was subjected to distillation under reduced pressure to leave 7.5 g (100%) of 3,4-dihydro-4-bromo-1-(2,4,6-triisopropylphenylsulfonyl) benz[cd] indol-5(1H)-one as a crude crystalline product. Then, this crude crystalline product was dissolved in a mixture of N,N-dimethylformamide (140 ml) and acetic acid (2 ml). To the solution was gradually added, while stirring at -28 °C, an aqueous solution (14 ml) of sodium azide (1.86 g, 28.6 mmol.) to raise the temperature of the reaction system up to near 0 °C. After completion of the reaction, the reaction mixture was added to ice-water (300 ml). Crystals then precipitated were collected by filtration and dissolved in ethyl acetate. The solution was shaken to allow two layers to be formed. The organic layer was washed with an aqueous saline solution, which was then dried (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure to

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leave 5.67 g (100%) of the title compound as a crude crystalline product. A portion of the product was recrystallized from ethyl acetate to give a product, m.p.129-132 °C.

¹H-NMR(CDCl₃, 200 MHz) δ: 1.12(d, J=7Hz, 12H), 1.28(d, J=7Hz, 6H), 2.94(t, J=7Hz, 1H), 3.16(ddd, J=16, 11, 2 Hz, 1H), 3.48(dd, J=16, 7Hz, 1H), 4.18(hep, J=7Hz, 2H), 4.49(dd, J=11, 7Hz, 1H), 7.21(s, 2H), 7.31(s, 1H), 7.40-(t, J=8Hz, 1H), 7.76(d, J=8Hz, 1Hx2) ppm. IR(KBr): 2970, 2110, 1695, 1600 cm⁻¹. Anal: Calcd for C₂₅H₃₀N₄O₃·0.5H₂O; C 64.04%, H 6.41%, N 11.49% Found; C 64.55%, H 6.27%, N 11.20%.

Reference Example 12

4-Acetylamino-3,4-dihydro-1-(2,4,6-triisopropylphenyl sulfonyl)benz[cd]indol-5(1H)-one

A solution of 4-azido-3,4-dihydro-1-(2,4,6-triisopropylphenylsulfonyl)benz[cd]indol-5(1H)-one (5.36 g, 11.2 mmol), acetic anhydride (2.29 g, 22.4 mmol) and 10% Pd-C (1.7 g in THF (60 ml) was stirred for 4 hours at 4 hours under 4 atmospheric pressure under hydrogen atmosphere. After completion of the reaction, the catalyst was filtered off. The filtrate was concentrated under reduced pressure. To the concentrate was added isopropyl ether to give 5.03 g (91%) of the title compound.

m.p.: 159-162 °C. ¹H-NMR(CDCl₃, 200 MHz) δ: 1.07, 1.10, 1.26(d, J=7Hz, 6Hx3), 2.13(s, 3H), 2.85(ddd, J=15, 12, 2 Hz, 1H), 2.92*hep, J=7Hz, 1H), 3.95(dd, J=15, 7Hz, 1H), 4.16(hep, J=7 Hz, 2H), 4.85(ddd, J=12, 7, 6Hz, 1H), 6.71(d, J=6Hz, 1H), 7.19(s, 2H), 7.23(d, J=2Hz, 1H), 7.38(t, J=8Hz, 1H), 7.68, 7.76(d, J=8Hz, 1H) ppm. IR(KBr); 3420, 3380, 2960, 1690, 1875, 1600 cm⁻¹.

Anal: Calcd. for C ₂₈ H ₃₄ N ₂ O ₄	C 67.99%,	H 6.93%,	N 5.66%
Found;	C 67.72%,	H 7.07%,	N 5.59%.

Reference Example 13

4-Trifluoroacetyl-3,4-dihydro-1-(2,4,6-triisopropylphenyl sulfonyl)benz[cd]indol-5(1H)-one

In 100 ml of ethanol was suspended 1.00 g (2.09 mmol) of 4-azido-3,4-dihydro-1-(2,4,6-triisopropylphenylsulfonyl)benz[cd]indol-5(1H)-one, to which was added 1.0 ml of conc. HCl. The mixture was warmed to make a solution, to which was added 0.10 g of 10% Pd-C (hydrous), followed by hydrogenation for one hour at room temperature. The reaction mixture was subjected to filtration to remove the catalyst. The filtrate was washed with ethanol and concentrated under reduced pressure. To the concentrate was added ether, which was concentrated twice, followed by crystallization from ether. The crystalline product was collected by filtration, washed with ether and dried under reduced pressure to afford 0.87 g of a 4-amino derivative. The yield was 85%.

0.40 g (0.818 mmol) of the 4-amino derivative obtained as above was dissolved in 6 ml of methylene chloride. To the solution was added, under ice-cooling, 0.14 ml (1.2 equivalent) of anhydrous trifluoroacetic acid and 0.27 ml of triethylamine. The mixture was stirred for 15 minutes, which was then concentrated under reduced pressure. To the concentrate were added ethyl acetate and 1N HCl, and the mixture was shaken to allow two layers to be formed. The organic layer was dried over anhydrous magnesium sulfate, which was concentrated under reduced pressure. The concentrate was purified by means of a silica gel column chromatography (elution with hexane - ethyl acetate 9:1). Fractions were combined and concentrated, to which was added a small volume of hexane to cause crystallization. The crystalline product was dried under reduced pressure to afford 0.23 g of the title compound. The yield was 51%. m.p.:155-156 °C. ¹H-NMR(CDCl₃, 200 MHz) δ: 1.09, 1.12, 1.26(d, J=7Hz, 6Hx3), 2.93(hep, J=7Hz, 1H), 2.96(ddd, 15, 12, 2 Hz, 1H), 4.04(dd, J=15, 7 Hz, 1H), 4.15(hep, J=7Hz, 2H), 4.83(ddd, J=12, 7, 6 Hz, 1H), 7.21(s, 2H), 7.29(d, J=2Hz, 1H), 7.41(t, J=8Hz, 1H), 7.67(d, J=6Hz, 1H), 7.73, 7.8(d, J=8Hz, 1H) ppm. IR(KBr): 2950, 1730, 1685, 1600 cm⁻¹. Anal: calcd for C₂₈H₃₁F₃N₂O₄S; C 61.30%, H 5.70%, N 5.11% Found; C 61.24%, H 5.80%, N 4.95%.

Example 1-1

4-Acetylamino-3,4-dihydro-5-hydroxy-1-(2,4,6-triisopropylphenylsulfonyl)-5-vinyl-5H-benz[cd]indole

To a solution of 4-acetylamino-3,4-dihydro-1-(2,4,6-triisopropylphenylsulfonyl)-benz[cd]indol-5(1H)-one (3.0 g, 6.0 mmol) in THF (60 ml) was added a solution of 1M vinyl magnesium bromide in THF (8 equivalents relative ketone compound) at -40°C in argon streams. The mixture was warmed up to room temperature, which was stirred for 4 hours at the same temperature range. The reaction mixture was added to an ice-cooled saturated aqueous solution of NH₄Cl. The mixture was subjected to extraction with ethyl acetate. The extract was washed with an aqueous saline solution and dried (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (1:2) to give 0.75 g (24%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200 MHz) δ : 1.08(d, J=7Hz, 12H), 1.24(d, J=7Hz, 6H), 2.00(s, 3H), 2.6-3.0(m, 2H), 3.18-3.31(dd, J=16.5Hz, 1H), 4.15(hept, J=7Hz, 2H), 4.51(dt, J=7, 4Hz, 1H), 5.31(dd, J=17, 10Hz, 1H), 5.50-5.73(dd, J=17, 10Hz, 1H), 7.1-7.4(m, 6H) ppm. IR(KBr): 3390, 2960, 1657 cm⁻¹. Anal: Calcd for C₃₀H₃₈N₂O₄S.H₂O; C 66.64%, H 7.4%, N 5.18% Found; C 66.35%, H 7.2%, N 4.7%.

Example 1-2

3,4-Dihydro-4-ethylamino-5-hydroxy-1-(2,4,6-triisopropylphenylsulfonyl)-5-vinyl-5H-benz[cd]indole

LAH (5 equivalents) was allowed to react with a THF solution (14.3 ml) of 4-acetylamino-3,4-dihydro-5-hydroxy-1-(2,4,6-triisopropylphenylsulfonyl)-5-vinyl-5H-benz[cd]indole (750 mg, 1.43 mmol) and was refluxed for 3 hours to complete the reaction. Ethanol was added to the reaction mixture to inactivate the excess amount of LAH and was added aqueous solution of Na₂SO₄. The reaction mixture was subjected to extraction with ethyl acetate. The extract was washed with an aqueous saline solution and, then, dried (anhydrous Na₂SO₄) and concentrated under reduced pressure. The residue thus obtained was subjected to a silica gel column chromatography, eluting with ethyl acetate to afford 275 mg (38%) of the title compound as an oily product.

¹H-NMR(CDCl₃, 200 MHz) δ : 1.11, 1.12(d, J=7Hz, 6Hx2), 1.25(d, J=7Hz, 6H), 2.4-3.1(m, 5H), 3.30-3.48(dd, J=15.4Hz, 1H), 4.18(hept, J=7Hz, 2H), 5.0-5.3(m, 2H), 5.92(dd, J=17, 11Hz, 1H), 7.1-7.4(m, 6H) ppm.

Example 1-3

Trans-4,6,6a,7,9,10a-hexahydro-7-ethyl-4-(2,4,6-triisopropylphenylsulfonyl)-10a-vinyl-indolo[3,4-gh][1,4]-benzoxazin-8-one

To a solution of 3,4-dihydro-4-ethylamino-5-hydroxy-1-(2,4,6-triisopropylphenylsulfonyl)-5-vinyl-5H-benz[cd]indole (270 mg, 0.53 mmol) in ethyl acetate (5.3 ml) was added a saturated aqueous solution of sodium carbonate (5.3 ml). To the mixture was added, while vigorously stirring, chloroacetyl chloride (1.2 equivalent). After completion of the reaction, the organic layer was separated and washed with an aqueous saline solution, followed by drying (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure to leave the residue (260 mg). To a solution of the chloroacetyl compound in THF-acetonitrile (5:1, 5ml) was added, while stirring under ice-cooling, 60% sodium hydride (2 equivalents), and the mixture was stirred for 2-3 hours at room temperature. After completion of the reaction, the reaction mixture was added to a saturated aqueous solution of NH₄Cl cooled with ice, followed by extraction with ethyl acetate. The extract solution was dried (anhydrous Na₂SO₄), then the solvent was distilled off. The residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (3:1) to afford 208 mg (86%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200MHz) δ : 1.07(d, J=7Hz, 12H), 1.24(d, J=7Hz, 6H), 2.7-3.0(m, 2H), 3.35(m, 2H), 3.9-4.2(m, 4H), 4.40, 4.48 (d, J=17Hz, 1Hx2), 4.91(d, J=18Hz, 1H), 5.36(d, J=11Hz, 1H), 6.24(dd, J=18, 11Hz, 1H), 7.1-7.4(m, 6H) ppm. IR(KBr): 2962, 1662 cm⁻¹. Anal: Calcd. for C₃₂H₄₀N₂O₄S.0.5H₂O; C 68.91%, H 7.41%, N 5.02% Found; C 68.86%, H 7.45%, N 4.69%.

Example 1-4

Trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-4-(2,4,6-triisopropylphenylsulfonyl)-10a-vinyl-7H-indolo[3,4-gh] [1.4]-benzoxazine

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To a solution of trans-4,6,6a,7,9,10a-hexahydro-7-ethyl-4-(2,4,6-triisopropylphenylsulfonyl)-10a-vinyl-indolo[3,4-gh][1.4]benzoxazin-8-one (200 mg, 0.36 mmol) in THF (3.6 ml) was added LAH (3 equivalents) while stirring under ice-cooling, and the mixture was heated for 3 hours under reflux. After completion of the reaction, ethanol was added to the reaction mixture to inactivate the excess amount of LAH, followed by addition to an ice-cooled saturated aqueous solution of Na₂SO₄. The mixture was subjected to extraction with ethyl acetate. The extract solution was washed with an aqueous saline solution and, then, dried (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure, and the residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (17:3) to afford 103 mg (54%) of the title compound as an amorphous crystalline product. ¹H-NMR(CDCl₃, 200MHz) δ: 1.06, 1.07 (d, J=7Hz, 6Hx2), 1.23(d, J=7Hz, 6H), 2.5-3.0(m, 7H), 3.21(dd, J=15, 4Hz, 1H), 3.80(dd, J=11, 3Hz, 1H), 4.1-4.4(m, 3H), 4.76(dd, J=18, 2Hz, 1H), 5.23(dd, J=11, 2Hz, 1H), 6.62(dd, J=18, 11Hz, 1H), 7.1-7.3(m, 6H) ppm. IR(KBr): 2975 cm⁻¹. Anal: Calcd. for C₃₂H₄₂N₂O₃S.0.5H₂O; C 70.68%, H 7.97%, N 5.15% Found; C 71.07%, H 7.85%, N 5.48%.

20 Example 1-5

Trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-10a-vinyl-7H-indolo[3,4-gh][1.4]benzoxazine

To a solution of trans-4,6,6a,8,9,10a-hexahydro-7-ethyl 1-4-(2,4,6-triisopropylphenylsulfonyl)-10a-vinyl-7H-indolo[3,4-gh] [1.4]benzoxazine (100 mg, 0.18 mmol) in THF (5.4 ml) was added, at room temperature, a solution of 0.5M naphthalene sodium in THF (10 equivalents) in argon streams. The mixture was stirred for 10 minutes at room temperature, which was then added to an ice-cooled saturated aqueous solution of NH₄Cl, followed by extraction with ethyl acetate. The organic layer was washed with an aqueous saline solution and dried (Na₂SO₄). The solvent was distilled off under reduced pressure, and the residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (1:1) to afford 33 mg (68%) of the title compound as an amorphous crystalline product. ¹H-NMR(CDCl₃, 200MHz) δ: 1.05(t, J=7Hz, 3H), 2.6-3.1(m, 6H), 3.31(dd, J=15, 4Hz, 1H), 3.83(ddd, J=11.4, 1Hz, 1H), 4.32(dt, J=11, 3Hz, 1H), 5.00(dd, J=17, 2Hz, 1H), 5.27(dd, J=11, 2Hz, 1H), 6.59(dd, J=17, 11Hz, 1H), 6.88(t, J=2Hz, 1H), 7.1-7.3(m, 3H), 7.8-8.0(br, 1H) ppm. IR(KBr): 3410, 2960 cm⁻¹. Anal: Calcd. for C₁₇H₂₀N₂O.0.9H₂O; C 71.75%, H 7.72%, N 9.84% Found; C 71.68%, H 7.25%, N 9.58%.

Example 2-1

4-Acetylamino-3,4-dihydro-5-hydroxy-5-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-5H-benz[cd]indole

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To a solution of 4-acetylamino-3,4-dihydro-1-(2,4,6-triisopropylphenylsulfonyl)-benz[cd]indol-5-one (3.57 g, 7.0 mmol) in THF (70 ml) was added, at 40 °C in argon streams, a solution of 1M methyl magnesium bromide in THF (8 equivalents relative to ketone compound). The mixture was warmed up to room temperature, and stirred for 4 hours. The reaction mixture was added to an ice-cooled saturated aqueous solution of NH₄Cl, which was subjected to extraction with ethyl acetate. The extract was washed with an aqueous saline solution and, then, dried (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with n-hexane-ethyl acetate, (1:2) to afford 1.27 g (36%) of the title compound as an amorphous crystalline product. ¹H-NMR(CDCl₃, 200MHz) δ: 1.09(d, J=7Hz, 12H), 1.25(d, J=7Hz, 6H), 1.95(s, 3H), 2.7-3.0(m, 2H), 3.24-(dd, J=16, 5Hz, 1H), 4.14(hept, J=7Hz, 2H), 4.54(m, 1H), 5.3-5.5(br, 1H), 7.1-7.4(m, 6H) ppm. IR(KBr): 3390, 2965, 1660 cm⁻¹. Anal: Calcd. for C₂₉H₃₈N₂O₄S.H₂O: C 65.88%, H 7.63%, N 5.30% Found; C 66.13%, H 7.28%, N 5.69%.

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Example 2-2

3,4-Dihydro-4-ethylamino-5-hydroxy-6-methoxy-5-methyl-5H-benz[cd]indole

- 5 In substantially the same manner as in Example 1-2, 4-acetylamino-3,4-dihydro-5-hydroxy-6-methoxy-5-methyl-1-p-toluenesulfonyl-5H-benz[cd]indole (2.0 g, 4.85 mmol) was subjected to reduction. The residue thus obtained was subjected to a silica gel column chromatography, eluting with ethyl acetate - methanol, (20:1) to afford 953 mg (76%) of the title compound as an amorphous crystalline product, m.p.185-187 °C (decomposed: as fumarate).
- 10 ¹H-NMR(CDCl₃, 200MHz) δ: 1.22(t, J=7Hz, 3H), 1.53(s, 3H), 2.5-2.8(m, 2H), 2.98(dq, J=11, 7Hz, 1H), 3.94-(s, 3H), 6.8-7.2(m, 3H), 7.96(br s, 1H) ppm. Anal: Calcd. for C₁₉H₂₄N₂O₆.0.5C₄H₄O₄; C 64.13%, H 6.96%, N 8.80% Found; C 63.77%, H 7.17%, N 8.79% as fumarate

Example 2-3

- 15 Trans-4,6,6a,7,9,10a-hexahydro-7-ethyl-1-methoxy-10a-methyl-indolo[3,4-gh][1.4]benzoxazin-8-one

- To a solution of 3,4-dihydro-4-ethylamino-5-hydroxy-6-methoxy-5-methyl-5H-benz[cd]indole (646 mg, 2.48 mmol) in ethyl acetate (25 ml) was added a saturated aqueous solution of sodium carbonate (24 ml). To the mixture was added, while stirring vigorously, chloroacetyl chloride (1.2 equivalent). After completion of the reaction, the organic layer was separated and washed with an aqueous saline solution, followed by drying (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure. The residue was cyclized and, then, purified by means of a silica gel column chromatography (eluting with n-hexane - ethyl acetate (2:7)) to afford 517 mg (72%) of the title compound as an amorphous crystalline product. ¹H-NMR(CDCl₃, 200MHz) δ: 1.21(t, d=7Hz, 3H), 1.57(s, 3H), 2.76(ddd, J=14, 13, 2 Hz, 1H), 3.38(m, 2H), 3.88(s, 3H), 4.0-4.3-(m, 2H), 4.44, 4.55 (d, J=17Hz, 1Hx2), 6.9-7.3(m, 3H), 7.94(br s, 1H) ppm. IR(KBr): 3348, 2988, 1645 cm⁻¹. Anal: Calcd for C₁₇H₂₀N₂O₃.1.5H₂O; C 62.37%, H 7.08%, N 8.56% Found; C 62.04%, H 6.30%, N 8.45%.
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Example 2-4

- 30 Trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-1-methoxy-10a-methyl-7H-indole[3,4-gh][1.4]benzoxazine

- Trans-4,6,6a,7,9,10a-hexahydro-7-ethyl-1-methoxy-10a-methyl-indolo[3,4-gh][1.4]benzoxazin-8-one (500 mg, 1.67 mmol) was subjected to reduction, which was recrystallized from chloroform - n-hexane to afford the title compound (320 mg, 67%, m.p.215-217 °C(dec.)). ¹H-NMR(CDCl₃, 200MHz) δ: 1.05(t, d=7Hz, 3H), 1.59(s, 3H), 2.4-3.0(m, 6H), 3.24(dd, J=17, 4Hz, 1H), 3.87-(s, 3H), 4.22(dt, J=12, 3Hz, 1H), 6.8-7.2(m, 3H), 7.78(br s, 1H) ppm. IR(KBr): 3320, 2959, 2803 cm⁻¹. Anal: Calcd for C₁₇H₂₂N₂O₂.0.3H₂O; C 69.97%, H 7.81%, N 9.60% Found; C 69.99%, H 7.60%, N 9.48%.
- 35

40 Example 3-1

3,4-Dihydro-5-hydroxy-5-methyl-4-trifluoroacetyl-amino-1-(2,4,6-triisopropylphenylsulfonyl)-5H-benz[cd]indole

- Methyl magnesium bromide (1M THF solution 8 equivalents) was allowed to react with a solution of 3,4-dihydro-4-trifluoroacetyl-amino-1-(2,4,6-triisopropyl phenylsulfonyl)-5H-benz[cd]indol-5-one (1.5 g, 1.43 mmol) in THF (14.3 ml). The reaction mixture was processed in a conventional manner. The residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (4:1) to afford 1.3 g (84%) of the title compound as an amorphous crystalline product.
- 50 ¹H-NMR(CDCl₃, 200MHz) δ: 1.08(d, J=7Hz, 12H), 1.25(d, J=7Hz, 6H), 1.59(s, 3H), 2.8-3.0(m, 2H), 3.40-(dd, J=18, 6Hz, 1H), 4.15(hept, J=7Hz, 2H), 4.55(m, 1H), 6.0-6.2(br, 1H), 7.2-7.5(m, 6H) ppm. IR(KBr): 3413, 2960, 1720 cm⁻¹. Anal: Calcd for C₂₉H₃₅N₂O₄ SF₃.0.5H₂O; C 60.72%, H 6.33%, N 4.88% Found; C 60.59%, H 6.18%, 4.55%.
- 55

Example 3-2

4-Amino-3,4-dihydro-5-hydroxy-5-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-5H-benz[cd]indole

To an ethanol solution (9 ml) of 3,4-dihydro-5-hydroxy-5-methyl-4-trifluoroacetyl-1-(2,4,6-triisopropylphenylsulfonyl)-5H-benz[cd]indole (640 mg, 0.91 mmol) was added 1N sodium hydroxide (2 equivalents). The mixture was heated for one hour under reflux. After completion of the reaction, the reaction mixture was concentrated under reduced pressure. The concentrate was shaken with chloroform-water, which was left standing to form two layers. The organic layer was separated and washed with an aqueous saline solution, followed by drying (anhydrous Na_2SO_4). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with ethyl acetate-methanol:triethylamine (50:5:1) to afford 290 mg (68%) of the title compound as an amorphous crystalline product.

$^1\text{H-NMR}(\text{CDCl}_3, 200\text{MHz})$ δ : 1.07(d, J = 7Hz, 12H), 1.24(d, J = 7Hz, 6H), 1.37(s, 3H), 2.89(hep, J = 7Hz, 1H), 3.1-3.4(m, 2H), 4.15(hep, J = 7Hz, 2H), 7.1-7.4(m, 2H) ppm. SIMS: $\text{MH}^+ = 469$.

Example 3-3

Trans-4,6,6a,7,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-indolo[3,4-gh][1.4] benzoxazin-8-one

To a solution of 4-amino-3,4-dihydro-5-hydroxy-5-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-5H-benz[cd] indole (234 mg, 0.5 mmol) in ethyl acetate (5 ml) was added a saturated aqueous solution of sodium carbonate (5 ml). To the mixture was added, while stirring vigorously, chloroacetyl chloride (1.2 equivalent). After completion of the reaction, the organic layer was separated and washed with an aqueous saline solution, followed by drying (anhydrous Na_2SO_4). The solvent was distilled off under reduced pressure. The residue was subjected to cyclization substantially in accordance with Example 1-3. The residue thus obtained was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (1:1) to afford 247 mg (99%) of the title compound as a crude crystalline product. A portion of the crystals was recrystallized (n-hexane - ethyl acetate) to give a pure product (m.p. 125-127 deg.)

$^1\text{H-NMR}(\text{CDCl}_3, 200\text{MHz})$ δ : 1.07, 1.08 (d, J = 7Hz, 6Hx2), 1.24(d, J = 7Hz, 6H), 1.45(s, 3H), 2.6-3.1(m, 3H), 4.00-(dd, J = 14, 6Hz, 1H), 4.15(hep, J = 7Hz, 2H), 4.47(s, 2H), 6.48(br s, 1H), 7.1-7.4(m, 6H) ppm. IR(KBr): 2960, 1684 cm^{-1} . SIMS: $\text{MH}^+ = 509$.

Example 3-4

Trans-4,6,6a,8,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-7H-indolo[3,4-gh][1.4] benzoxazine

To a solution of trans-4,6,6a,7,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-indolo[3,4-gh][1.4]benzoxazin-8-one (220 mg, 0.43 mmol) in THF (10 ml) was added, while stirring under ice-cooling, LAH (3 equivalents). The mixture was heated for 3 hours under reflux. After completion of the reaction, ethanol was added to the reaction mixture to inactivate the excess amount of LAH. The reaction mixture was added to an ice-cooled saturated aqueous solution of Na_2SO_4 , followed by extraction with ethyl acetate. The extract solution was washed with an aqueous saline solution, followed by drying (anhydrous Na_2SO_4). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (1:5), to afford 106 mg (50%) of the title compound as a syrupy product.

$^1\text{H-NMR}(\text{CDCl}_3, 200 \text{ MHz})$ δ : 1.07(d, J = 7Hz, 12H), 1.23(d, J = 7Hz, 6H), 1.44(s, 3H), 2.58(ddd, J = 14, 12, 2 Hz, 1H), 2.7-3.2(m, 5H), 3.79(dd, J = 12, 3Hz, 1H), 4.0-4.3(m, 3H), 7.1-7.3(m, 6H) ppm.

Example 3-5

Trans-4,6,6a,8,9,10a-hexahydro-10a-methyl-7H-indolo[3,4-gh][1.4]benzoxazine

Lithium aluminum hydride (5 equivalents) was allowed to react with a THF solution (20 ml) of trans-4,6,6a,8,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropyl phenylsulfonyl)-7H-indolo[3,4-gh][1.4] benzoxazin-8-one (1.2 g, 2.36 mmol). The residue thus obtained was subjected to a silica gel column chromatography, eluting with ethyl acetate, to afford 100 mg (15%) of the title compound.

¹H-NMR(CDCl₃, 200MHz) δ: 1.51(s,3H), 2.66(ddd,J = 15, 12, 2 Hz, 1H), 2.8-3.3(m,4H), 3.82(dd, J = 11, 3 Hz, 1H), 4.13(dt,J = 12, 3Hz, 1H), 6.87(br s, 1H), 7.1-7.2(m,3H), 8.0-8.1(br,1H) ppm. IR(KBr): 3410, 2935 cm⁻¹. Anal: Calcd for C₁₄H₁₆N₂O.1.0H₂O; C 68.27%, H 7.37%, N 11.37% Found; C 68.12%, H 7.01%, N 11.10%.

5 Example 4

4,6,6a,8,9,10a-Hexahydro-10a-methyl-7-(4-phthalimidobutyl)-7H-indolo[3,4-gh][1.4] benzoxazine

To a solution of 4,6,6a,8,9,10a-hexahydro-10a-methylindolo[3,4-gh][1.4]benzoxazine (100 mg, 0.44 mmol) in DMF (5 ml) were added potassium carbonate (303 mg, 2.2 mmol), N-(4-bromobutyl)phthalimido (124 mg, 0.44 mmol) and sodium iodide (0.66 mg, 0.44 mmol). The mixture was heated at 60 °C for 24 hours. After completion of the reaction, the reaction mixture was added to ice water - ethyl acetate. The organic layer was separated, washed with a saturated aqueous saline solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (3:2), to afford 130 mg (69%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200 MHz) δ: 1.52(s,3H), 1.5-1.9(m,4H), 2.3-3.0(m,6H), 3.28(dd,J = 14, 4Hz,1H), 3.75-(t,J = 7Hz,2H), 4.18(dt,J = 12, 3Hz,1H), 6.90(br s, 1H), 7.1-7.2(m,3H), 7.7-7.8(m,2H), 7.8-7.9(m,2H), 7.96(br s, 1H) ppm. IR(KBr): 3411, 2945, 1710 cm⁻¹. Anal: Calcd for C₂₆H₂₇N₃O₃.1.0H₂O; C 69.78%, N 9.39% Found; C 69.61%, H 6.18%, N 8.90%.

Example 5

4,6,6a,8,9,10a-Hexahydro-10a-methyl-7-(2-phenoxyethyl)-7H-indolo[3,4-gh][1.4]benzoxazine

To a solution of 4,6,6a,8,9,10a-hexahydro-10a-methyl-7H-indolo[3,4-gh][1.4]benzoxazine (85 mg, 0.37 mmol) in DMF (5 ml) were added potassium carbonate (255 mg, 1.9 mmol) and 1-iodo-2-phenoxyethane (99 mg, 0.4 mmol). The mixture was heated at 60 °C for 24 hours. After completion of the reaction, the reaction mixture was added to a mixture of ice water and ethyl acetate. The organic layer was separated, washed with a saturated aqueous saline solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure, and the residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (2:1), to afford 59 mg (46%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200MHz) δ: 1.54(s,3H), 2.5-3.0(m,5H), 3.2-3.4(m,2H), 3.82(dd,J = 12,3 Hz,1H), 4.1-4.3(m,3H), 6.9-7.9(m,9H), 7.95(br s,1H) ppm. IR(KBr): 3412, 2947, 1710 cm⁻¹. Anal: Calcd for C₂₂H₂₄N₂O₂.0.5H₂O; C 73.92%, H 7.05%, N 7.84% Found; C 73.44%, H 6.82%, N 7.10%.

Example 6-1

40 Trans-7-chloroacetyl-4,6,6a,8,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-7H-indolo[3,4-gh][1.4] benzoxazine

To a solution of trans-4,6,6a,8,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-7H-indolo[3,4-gh][1.4] benzoxazine (106 mg, 0.21 mmol) in ethyl acetate (10 ml) was added a saturated aqueous solution of sodium carbonate (2 ml). To the mixture was added, while stirring vigorously, chloroacetyl chloride (1.2 equivalent). After completion of the reaction, the organic layer was separated, washed with an aqueous saline solution and dried (anhydrous Na₂SO₄). The solvent was distilled off under reduced pressure, and the residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (4:1) to afford 109 mg (89%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200MHz) δ: 1.08, 1.085(d,J = 7Hz, 6Hx2), 1.24(d,J = 7Hz,6H), 2.89(hep,J = 7Hz,1H), 3.45-(dd,J = 17, 4Hz,1H), 3.68(dd,J = 12, 4Hz,1H), 3.8-4.2(m,7H), 7.1-7.4(m,6H) ppm. IR(KBr): 2965, 1663 cm⁻¹. Anal: Calcd for C₃₁H₃₉N₂O₄SCl; C 65.19%, H 6.88%, N 4.90% Found; C 64.78%, H 6.84%, N 4.82%.

Example 6-2

Trans-7-[2-(4-benzylpiperazin-1-yl)-1-oxoethyl]-4,6,6a,8,9,10a-hexahydro-10a-methyl-4-(2,4,6-triisopropylphenylsulfonyl)-7H-indolo[3,4-gh][1.4] benzoxazine

5

To a solution of trans-7-chloroacetyl-4,6,6a,8,9,10a-hexahydro-10a-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-7H-indolo[3,4-gh][1.4] benzoxazine (77 mg, 0.14 mmol) in DMF (2.5 ml) were added potassium carbonate (82 mg, 0.6 mmol) and benzyl piperazine (0.025 ml, 0.14 ml). The mixture was stirred for 8 hours at room temperature. After completion of the reaction, a mixture of ice-water and ethyl acetate was added to the reaction mixture. The organic layer was separated, washed with a saturated aqueous saline solution and dried over anhydrous sodium sulfate. Under reduced pressure, the solvent was distilled off to leave 81 mg (85%) of the title compound as a crystalline product. A portion of the crystals was recrystallized from ether - n-hexane (m.p. 82-84 deg.)

¹H-NMR(CDCl₃, 200MHz) δ: 1.07, 1.08 (d, J = 7Hz, 6Hx2), 1.23(d, J = 7Hz, 6H), 1.50(s, 3H), 2.3-2.7(m, 8H), 3.10, 3.29 (d, J = 14 Hz, 1Hx2), 3.52(s, 2H), 3.61(dd, J = 12, 4 Hz, 1H), 3.8-4.2(m, 3H), 7.1-7.4(m, 11H) ppm. IR(KBr): 2960, 1652 cm⁻¹. Anal: Calcd for C₄₂H₅₄N₄O₄·2.0H₂O; C 67.53%, H 7.83%, N 7.50% Found; C 67.47%, H 7.31%, N 7.27%.

Example 6-3

20

Trans-7-[2-(4-benzylpiperazin-1-yl)-ethyl]-4,6,6a,8,9,10a-hexahydro-10a-methyl-7H-indolo[3,4-gh][1.4]-benzoxazine

7-[2-(4-benzylpiperazin-1-yl)-1-oxoethyl]-4,6,6a,8,9,10a-hexahydro-10a-methyl-4-(2,4,6-triisopropylphenylsulfonyl)-7H-indolo[3,4-gh][1.4]benzoxazine (71 mg, 0.1 mmole) was subjected to LAH reduction. The compound thus reduced was subjected to a silica gel column chromatography, eluting with ethyl acetate:methanol (20:1), to afford 29 mg (67%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200MHz) δ: 1.51(s, 3H), 2.4-3.1(m, 16H), 3.30(dd, J = 14, 4Hz, 1H), 3.52(s, 2H), 3.81(dd, J = 12, 3Hz, 1H), 4.18(dt, J = 12, 3Hz, 1H), 6.88(br s, 1H), 7.1-7.4(m, 8H), 7.97(br s, 1H) ppm. IR(KBr): 3420, 2940, 2810 cm⁻¹. Anal: Calcd for C₂₇H₃₄N₄O·2.0H₂O; C 69.50%, H 8.21%, N 12.0% Found; C 69.46%, H 7.53%, N 11.8%.

Example 7-1

4-Acetylamino-3,4-dihydro-5-hydroxy-6-methoxy-5-phenyl-1-p-toluenesulfonyl-5H-benz[cd]indole

In substantially the same manner as Example 1-1, phenyl magnesium bromide (2M THF solution) was allowed to react with 4-acetylamino-3,4-dihydro-6-methoxy-1-p-toluenesulfonylbenz[cd] indol-(1H)-5-one (2.0 g, 4.85 mmol). The residue thus obtained was subjected to a silica gel column chromatography, eluting with ethyl acetate, to afford 1.1 g (46.2%) of the title compound as an amorphous crystalline product.

¹H-NMR(CDCl₃, 200MHz) δ: 1.93(s, 3H), 2.35(ddd, J = 16, 13, 2 Hz, 1H), 2.38(s, 3H), 2.98(dd, J = 15, 5Hz, 1H), 3.71(s, 3H), 4.72(m, 1H), 4.94(s, 1H), 5.14(d, J = 10Hz, 1H), 6.92(d, J = 9Hz, 1H), 7.1-7.9(m, 11H) ppm. IR(KBr): 3510, 3424, 1661 cm⁻¹. Anal: Calcd for C₂₇H₂₆N₂O₅S; C 66.10%, H 5.3%, N 5.71% Found; C 65.93%, H 5.60%, N 5.61%.

Example 7-2

Trans-7-ethyl-4,6,6a,7,9,10a-hexahydro-1-methoxy-10a-phenyl-indolo[3,4-gh][1.4]benzoxazin-8-one

50

4-Acetylamino 3,4-dihydro-5-hydroxy-6-methoxy-1-p-toluenesulfonyl-5H-benz[cd]indole (600 mg, 1.86 mmol) was likewise cyclized by processing with chloroacetyl chloride according to the method of Example 2-3 to give crude crystals (692 mg). A portion of the crystals was recrystallized (chloroform-n-hexane) to afford a pure product (m.p 278-280 dec.).

¹H-NMR(CDCl₃, 200MHz) δ: 0.81(t, J = 7Hz, 3H), 3.69(s, 3H), 6.95(d, J = 9Hz, 1H), 7.2-7.4(m, 7H), 8.00(br s, 1H) ppm. IR(KBr): 1655 cm⁻¹.

Example 7-3

Trans-7-ethyl-1-methoxy-10a-phenyl-4,6,6a,10a-tetrahydro-7H-indolo[3,4-gh][1,4]benzoxazine

- 6 Trans-7-ethyl-4,6,6a,8,9,10a-hexahydro-1-methoxy-10a-phenyl-7H-indolo[3,4-gh][1,4]benzoxazin-8-one (664 mg, 1.83 mmol) was reduced with LAH. The residue thus obtained was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (3:1), to afford 74 mg (12%) of the title compound as a crude crystalline product (m.p.194-196 dec.)
 10 ¹H-NMR(CDCl₃, 200MHz) δ: 0.97(t, J = 7Hz, 3H), 2.7-3.1(m, 3H), 3.92(dd, J = 15, 5Hz, 1H), 3.61(s, 3H), 3.70-(dd, J = 13, 5Hz, 1H), 5.42, 6.37(d, J = 5Hz, 1Hx2), 6.88(d, J = 9Hz, 1H), 7.02(br s, 1H), 7.1-7.4(m, 7H), 7.91(br s, 1H) ppm. IR(KBr): 3355, 2967 cm⁻¹. Anal: Calcd for C₂₂H₂₂N₂O₂·0.5H₂O; C 74.3%, H 6.52%, N 7.88% Found; C 73.91%, H 6.20%, N 7.73%

Example 8-1

- 15 Trans-4-acetylamino-3,4-dihydro-5-hydroxy-5-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-5(1H)benz[cd]indole

To a solution of 4-acetylamino-3,4-dihydro-1-(2,4,6-triisopropylphenylsulfonyl)-benz[cd]indol-5(1H)-one (6.0 g, 12.1 mmol) in THF (100 ml) was added at - 40 °C, in argon streams, a THF solution (97 ml) of 1M
 20 methyl magnesium bromide. The mixture was warmed up to room temperature and stirred for 4 hours at the same temperature range, which was added to an ice-cooled saturated aqueous solution of ammonium chloride. The mixture was subjected to extraction with ethyl acetate. The extract was washed with a saturated aqueous saline solution, which was dried (anhydrous sodium sulfate). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with n-
 25 hexane - ethyl acetate (1:5) to afford 3.6 g (58%) of the title compound as amorphous crystals.
¹H-NMR(CDCl₃, 200 MHz) δ : 1.09(d, J = 7Hz, 12H), 1.25(d, J = 7Hz, 6H), 1.95(s, 3H), 2.70-3.00(m, 2H), 3.24(dd, J = 16.5Hz, 1H), 4.14(hept, J = 7Hz, 2H), 4.54(m, 1H), 5.30-5.50(br, 1H), 7.10-7.40(m, 6H) ppm. IR(KBr): 3390, 2965, 1660cm⁻¹. Anal: Calcd for C₂₉H₃₈N₂O₄S·H₂O; C 65.88%, H 7.63%, N 5.30% Found; C 66.13%, H 7.28%, N 5.69%.

- 30 Example 8-2

Trans-3,4-dihydro-4-ethylamino-5-hydroxy-5-methyl-1-2,4,6-triisopropylphenylsulfonyl)-5(1H)-benz[cd]indole

- 35 To a solution of trans-4-acetylamino-3,4-dihydro-5-hydroxy-5-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-5(1H)-benz[cd]indole in THF (50 ml) was added LAH (5 equivalents). The mixture was heated for 2 hours under reflux. After completion of the reaction, ethanol was added to the reaction mixture to inactivate excess LAH, which was then added to an ice-cooled saturated aqueous solution of sodium sulfate. The mixture was subjected to extraction with ethyl acetate, and the extract was washed with a saturated aqueous saline
 40 solution, followed by drying (anhydrous sodium sulfate). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with ethyl acetate to afford 3.05 g (90%) of the title compound as an oily product.
¹H-NMR(CDCl₃, 200 MHz) δ : 1.07, 1.08(d, J = 7Hz, 6Hx2), 1.15(t, J = 7Hz, 3H), 1.24(d, J = 7Hz, 6H), 2.35-3.10-(m, 5H), 3.30(dd, J = 15.4Hz, 1H), 4.15(hept, J = 7Hz, 2H), 7.10-7.35(m, 6H) ppm.

- 45 Example 8-3

Trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-10a-methyl-7H-indolo[3,4-gh][1,4]benzoxazine

- 50 To a solution of trans-3,4-dihydro-4-ethylamino-5-hydroxy-5-methyl-1-(2,4,6-triisopropylphenylsulfonyl)-5(1H)-benz[cd]indole (3.0 g, 6.0 mmol) in ethyl acetate (60 ml) was added a saturated aqueous solution of sodium carbonate (60 ml). To the solution was added, while vigorously stirring, chloroacetyl chloride (1.2 equivalent). After completion of the reaction, the organic layer was taken and washed with an aqueous saline solution, followed by drying (anhydrous sodium sulfate). The solvent was distilled off under reduced
 55 pressure to leave a residue (3.4 g). To a solution of this chloroacetyl compound in THF-DMF (5:3, 80 ml) was added, while stirring under ice-cooling, 60% sodium hydride (2 equivalents). The mixture was stirred for one hour at room temperature. After completion of the reaction, the reaction mixture was added to an ice-cooled saturated aqueous solution of ammonium chloride. The mixture was subjected to extraction with

ethyl acetate. The extract was further dried (anhydrous sodium sulfate), then the solvent was distilled off under reduced pressure to leave a residue (2.8 g). To a solution of thus-obtained residue in THF (50 ml) was added, at room temperature, LAH (8 equivalents), and the mixture was heated for 12 hours under reflux. After completion of the reaction, ethanol was added to the reaction mixture. The mixture was added to an ice-cooled saturated aqueous solution of sodium sulfate, followed by extraction with ethyl acetate. The extract was washed with an aqueous saline solution and dried (anhydrous sodium sulfate). The solvent was distilled off, and the residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (1:1). The eluate was recrystallized from isopropyl ether - n-hexane to give 600 mg (39%) of the title compound, m.p.129-130 °C

¹H-NMR(CDCl₃, 200 MHz) δ : 1.04(t, J = 7Hz, 3H), 1.53(s, 3H), 2.45-3.05(m, 6H), 3.28(dd, J = 14, 4Hz, 1H), 3.84-(dd, J = 11, 3Hz, 1H), 4.21(dt, J = 12, 3Hz, 1H), 6.88(s, 1H), 7.10-7.25(m, 3H), 7.85-8.05(br, 1H) ppm. IR(KBr): 3275, 2942, 1615, 1602 cm⁻¹. Anal: Calcd for C₃₁H₄₀N₂O₅; C 74.97%, H 7.86%, N 10.93% Found; C 74.90%, H 7.94%, N 10.92%

Example 9-1

Trans-4,6,6a,7,9,10a-hexahydro-7-methyl-10a-methyl-1-methoxy-4-(2,4,6-triisopropylphenylsulfonyl)-indolo[3,4-g h][1,4]benzoxazin-8-one

To a solution of 3,4-dihydro-5-hydroxy-6-methoxy-5-methyl-4-trifluoroacetyl-amino-1-(2,4,6-triisopropylphenylsulfonyl)-5H-benz[cd]indole (m.p.169-171 °C, Anal: Calcd. for C₂₉H₃₃N₂O₅SF₃•0.2H₂; C 59.82%, H 5.78%, N 4.81% Found; C 59.59%, H 5.88%, N 4.92%) (3.4 g, 5.7 mmol.), which was synthesized in substantially the same manner as in Reference Example 13, in ethanol (55 ml) was added a 1 N aqueous solution of sodium hydroxide (14.3 ml). The mixture was stirred for 30 minutes at 60 °C. After completion of the reaction, the solvent was distilled off under reduced pressure. The residue was subjected to extraction with chloroform, and the extract was washed with an aqueous saline solution, followed by drying (anhydrous sodium sulfate). The solvent was distilled off under reduced pressure to leave an amine compound. Then, to a solution of this amine compound in ethyl acetate (50 ml) was added a saturated aqueous solution of sodium carbonate (50 ml). To the mixture was added, while vigorously stirring, chloroacetyl chloride (1.2 equivalent). After completion of the reaction, the organic layer was taken and washed with an aqueous saline solution, followed by drying (anhydrous sodium sulfate). The solvent was distilled off under reduced pressure to leave a residue (3.2 g). To a solution of this chloroacetyl compound (2.5 g, 4.35 mmol) in THF (40 ml) were added, while stirring under ice-cooling, 60% sodium hydride (3 equivalents) and methyl iodide (5 equivalents). The mixture was stirred for 5 days at room temperature. After completion of the reaction, the reaction mixture was added to an ice-cooled aqueous solution of ammonium chloride. The mixture was subjected to extraction with ethyl acetate. The extract was further dried (anhydrous sodium sulfate), then the solvent was distilled off to leave a residue. The residue was subjected to a silica gel column chromatography, eluting with n-hexane - ethyl acetate (1:1-2:3) to afford 1.9 g (80%) of the title compound as amorphous crystals, m.p.155-157 °C (n-hexane - ethyl acetate).

¹H-NMR(CDCl₃, 200 MHz) δ : 1.08, 1.09(d, J = 7Hz, 6Hx2), 1.25(d, J = 7Hz), 1.51(s, 3H), 3.13(s, 3H), 3.15-(dd, J = 12, 4Hz, 1H), 3.86(s, 3H), 3.95-4.25(m, 3H), 4.48(ABq, 2H), 6.85-7.30(m, 3H) ppm. IR(KBr): 2960, 1655, 1600, 1429 cm⁻¹. Anal: Calcd for C₃₁H₄₀N₂O₅S; C 67.36%, H 7.29%, N 5.07% Found; C 67.44%, H 7.40%, N 5.09%.

Example 9-2

Trans-4,6,6a,8,9,10a-hexahydro-1-methoxy-7-methyl-10a-methyl-7H-indolo[3,4-gh][1,4]benzoxazine

To a solution of trans-4,6,6a,7,9,10a-hexahydro-7-methyl-10a-methyl-1-methoxy-4-(2,4,6-triisopropylphenylsulfonyl)-indolo[3,4-gh][1,4]benzoxazin-8-one (1.9 g, 3.4 mmol) in THF (30 ml) was added, while stirring under ice-cooling, LAH (3 equivalents). The mixture was heated for 3 hours under reflux. After completion of the reaction, ethanol was added to the reaction mixture, which was then added to an ice-cooled saturated aqueous solution of sodium sulfate. The mixture was subjected to extraction with ethyl acetate. The extract was washed with an aqueous saline solution, which was then dried (anhydrous sodium sulfate). The solvent was distilled off under reduced pressure. The residue was subjected to a silica gel column chromatography, eluting with ethyl acetate to leave 700 mg of a residual substance. To a solution of this residual substance in THF (10 ml) was added, at room temperature in argon streams, a solution of 0.5M

naphthalene sodium in THF (10 equivalents). The mixture was stirred for 10 minutes at room temperature, which was added to an ice-cooled saturated aqueous solution of ammonium chloride, followed by extraction with ethyl acetate. The organic layer was washed with an aqueous saline solution, which was dried (anhydrous sodium sulfate). The solvent was distilled off under reduced pressure. The residue was

5 subjected to a silica gel column chromatography, eluting with ethyl acetate to afford 250 mg (27%) of the title compound as crystals, m.p. 200-202 °C (isopropyl ether - ethyl acetate).
¹H-NMR(CDCl₃, 200 MHz) δ : 1.62(s,3H), 2.38(s,3H), 2.40-2.60(m,3H), 2.77(dd,J = 11,2Hz,1H), 3.26(m,1H), 4.26(dt,J = 12,3Hz,1H), 6.89(d,J = 9Hz,1H), 7.15(d,J = 9Hz,1H), 7.85-8.00(br,1H) ppm. IR(KBr): 3304, 1502, 1431 cm⁻¹. Anal: Calcd for C₁₆H₂₀N₂O₂; C 70.56%, H 7.40%, N 10.29% Found; C 70.67%, H 7.51%, N 10.45%

Biological Test Example 1

Action of Substance-P for inhibiting biting response by its administration at subarachnoid space

15 Male Jcl:ICR mice of five-week old (10 animals per group) were employed. Dorsal cutaneous incisions were made under anesthesia at least one hour before the experiments. Substance-P (SP) was administered in an amount of 10 ng/5 μl/mouse at the subarachnoid space of the spinal cord. Then the test animals started to bite themselves at the abdomen alternately. The number of bitings was counted for one minute

20 immediately after the SP administration. The values thus obtained were made indexes of pseudo pain reflexes caused by SP.
 Test drugs were dissolved in a solution (Tween 80 : ethanol : physiological saline = 20:10:70) together with a given amount of SP. The solutions thus prepared were administered to test animals. In the case of oral administration, 5% suspension of gum arabica was employed, which was administered at 30 minutes

25 before the SP administration.
 The inhibition rate (%) of each sample was calculated by the following formula.

Inhibition rate (%) = (number of counts in the control group - number of counts in each sample)/(number of counts in the control group) x 100.

30 And, 50% inhibition dose (ID₅₀) was calculated from the primary linear regression line of dose-dependent curve.

Table 1

Antagonistic action to Substance-P (Dosage 10 μg/mouse)	
Compounds (Example No.)	Inhibition rate (%)
1	80
2	100
8	65

Biological Test Example 2

Acetic acid - writhing method

50 Male Slc:ICR mice of four-week old (10 animals per group) were employed. Test compounds were orally administered. Thirty minutes later, purified water containing 0.6% acetic acid was injected intraperitoneally (0.1 ml/10 g body weight). Writhing shown by the test animals during 20 minutes was observed on individual animals.

On each test animal, the inhibition rate against the average response times in the control group was

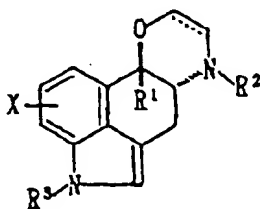
55 calculated. The test compound was administered as a 5% suspension of gum arabica.

Table 2

Analgesic action		
Compound (Example No.)	Inhibition rate (%)	Dose (mg/kg,p.o.)
2	99	25
8	61	12.5

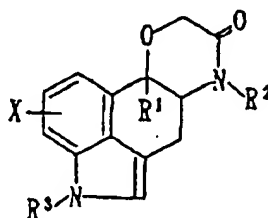
Claims

1. A compound of the formula



- wherein X is hydrogen, or a substituent; R¹ is an optionally substituted hydrocarbon residue; R² and R³ are respectively hydrogen or a substituent; and _____ shows a single bond or a double bond, or a pharmacologically acceptable acid addition salt thereof.
2. The compound of claim 1, wherein X is hydrogen, halogen or an optionally substituted hydrocarbon residue which may be bond to the ring through oxygen atom.
3. The compound according to claim 2, wherein X is hydrogen, lower alkyl, lower alkoxy or halogen.
4. The compound according to claim 3, wherein X is hydrogen or C₁₋₆ alkoxy.
5. The compound according to claim 1, wherein R¹ is lower alkyl, lower alkenyl, lower alkynyl or aryl.
6. The compound according to claim 5, wherein R¹ is C₁₋₆ alkyl, C₂₋₆ alkenyl or phenyl.
7. The compound of claim 1, wherein R² is hydrogen or an optionally substituted hydrocarbon residue.
8. The compound according to claim 7, wherein R² is hydrogen or lower alkyl which may be substituted by halogen, aryl, cycloalkyl, heterocyclic ring, aralkyl or acyl, and the aryl and heterocyclic ring may be bound to the lower alkyl through oxygen, sulfur or nitrogen atom.
9. The compound according to claim 8, wherein R² is hydrogen, C₁₋₆ alkyl or a group of the formula: -(CH₂)_nAR⁴ (wherein n is integer of 1 to 6; A is a bond or oxygen, sulfur or nitrogen atom; R⁴ is hydrogen, phenyl, 1-phthalimidyl, 4-benzylpiperazyl).
10. The compound of claim 1, wherein R³ is hydrogen, an optionally substituted hydrocarbon residue or a sulfonyl group.
11. The compound according to claim 10, wherein R³ is hydrogen, lower alkyl which may be substituted by halogen or arylsulfonyl which may be substituted by lower alkyl.
12. The compound according to claim 11, wherein R³ is hydrogen or sulfonyl which may be substituted by C₁₋₆ alkyl phenyl or tolyl.

13. The compound according to claim 1, wherein X is hydrogen or C₁₋₃ alkoxy; R¹ is C₁₋₃ alkyl, vinyl or phenyl; R² is hydrogen, C₁₋₆ alkyl or the group of the formula: -(CH₂)_nAR⁴; (wherein n is integer of 1 to 6; A is a bond or oxygen, sulfur or nitrogen atom; R⁴ is hydrogen, phenyl, 1-phthalimidyl, 4-benzylpiperazyl); R³ is hydrogen or 2,4,6-triisopropylphenylsulfonyl; and _____ is a single bond.
14. The compound according to anyone of claims 1 to 13, which is trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-10a-vinyl-7H-indolo[3,4-gh][1.4]benzoxazine.
15. The compound according to anyone of claims 1 to 13, which is trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-1-methoxy-10a-methyl-7H-indolo[3,4-gh][1.4]benzoxazine.
16. The compound according to anyone of claims 1 to 13, which is trans-4,6,6a,8,9,10a-hexahydro-7-ethyl-10a-methyl-7H-indolo[3,4-gh][1.4]benzoxazine fumalate.
17. A method of producing the compound according to claim 1, which comprises subjecting a compound of the formula:



wherein X, R¹, R² and R³ are of the same meaning as defined above, to reduction, and, when desired, by further subjecting the reduced compound to reduction, oxidation or/and substitution reaction.

18. A pharmaceutical composition which contains the compound according to anyone of claims 1 to 16, together with pharmaceutically acceptable carrier, vehicle and diluent.
19. The compound of anyone of claims 1 to 16, as a pharmaceutical active ingredient.
20. Use of the compound according to anyone of claims 1 to 16 for manufacturing a drug intended for prophylaxis or treatment of a mammal suffering from pain.



European Patent
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PARTIAL EUROPEAN SEARCH REPORT

Application Number

which under Rule 45 of the European Patent Convention EP 94 10 1551
shall be considered, for the purposes of subsequent
proceedings, as the European search report

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.5)
D,A	EP-A-0 037 784 (ROUSSEL-UCLAF) * claims 1,13 * ---	1,18	C07D498/06 A61K31/535 //(C07D498/06, 265:00,209:00)
D,A	US-A-4 238 486 (J. H. JONES) * claims 1,16,19 * ---	1,18	
A	CHEMICAL ABSTRACTS, vol. 105, no. 19, 10 November 1986, Columbus, Ohio, US; abstract no. 164465t, J. H. JONES ET AL 'Oxazine derivatives as dopamine antagonists' page 18 ; * abstract * & SPEC. PUBL. - R. SOC. CHEM. vol. 55 , 1986 pages 96 - 108 -----	1,18	
			TECHNICAL FIELDS SEARCHED (Int.Cl.5)
			C07D A61K
INCOMPLETE SEARCH			
<p>The Search Division considers that the present European patent application does not comply with the provisions of the European Patent Convention to such an extent that it is not possible to carry out a meaningful search into the state of the art on the basis of some of the claims</p> <p>Claims searched completely : Claims searched incompletely : Claims not searched : Reason for the limitation of the search:</p> <p>see sheet C</p>			
Place of search		Date of completion of the search	Examiner
THE HAGUE		24 May 1994	Voyiazoglou, D
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons ----- & : member of the same patent family, corresponding document	



EP 94 10 1551 -C-

INCOMPLETE SEARCH

Claims searched completely : 13-16
Claims searched incompletely : 1-12,17-20

Reason : The expressions "substituent" and "optionally substituted hydrocarbon residue" are not clear and concise (Article 84 EPC). Guided by the spirit of the application and the inventive concept as disclosed in the descriptive part of the present application the search has been based on the examples.